

Valentina Zampiga

List of Publications by Year in descending order

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papers

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1163117

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13
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957
citing authors

#	ARTICLE	IF	CITATIONS
1	Male Breast Cancer: Results of the Application of Multigene Panel Testing to an Italian Cohort of Patients. <i>Diagnostics</i> , 2020, 10, 269.	2.6	14
2	The Spectrum of FANCM Protein Truncating Variants in European Breast Cancer Cases. <i>Cancers</i> , 2020, 12, 292.	3.7	11
3	Insights into Genetic Susceptibility to Melanoma by Gene Panel Testing: Potential Pathogenic Variants in ACD, ATM, BAP1, and POT1. <i>Cancers</i> , 2020, 12, 1007.	3.7	19
4	Identification of a novel large EPCAM-MSH2 duplication, concurrently with LOHs in chromosome 20 and X, in a family with Lynch syndrome. <i>International Journal of Colorectal Disease</i> , 2019, 34, 1999-2002.	2.2	0
5	Multigene Panel Testing Increases the Number of Loci Associated with Gastric Cancer Predisposition. <i>Cancers</i> , 2019, 11, 1340.	3.7	19
6	Large scale multifactorial likelihood quantitative analysis of <i>BRCA1</i> and <i>BRCA2</i> variants: An ENIGMA resource to support clinical variant classification. <i>Human Mutation</i> , 2019, 40, 1557-1578.	2.5	102
7	Multiple-gene panel analysis in a case series of 255 women with hereditary breast and ovarian cancer. <i>Oncotarget</i> , 2017, 8, 47064-47075.	1.8	68
8	BRCA1 p.His1673del is a pathogenic mutation associated with a predominant ovarian cancer phenotype. <i>Oncotarget</i> , 2017, 8, 22640-22648.	1.8	10
9	Multiple Primary Tumors in a Family with Li-Fraumeni Syndrome with a TP53 Germline Mutation Identified by Next-Generation Sequencing. <i>International Journal of Biological Markers</i> , 2016, 31, 461-465.	1.8	2
10	First evidence of a large CHEK2 duplication involved in cancer predisposition in an Italian family with hereditary breast cancer. <i>BMC Cancer</i> , 2014, 14, 478.	2.6	12
11	Characterization of Molecular Alterations of BRCA1/2: Analysis and Interpretation Guidelines. <i>Current Women's Health Reviews</i> , 2012, 8, 4-11.	0.2	0
12	Results of a population-based screening for hereditary breast cancer in a region of North-Central Italy: contribution of BRCA1/2 germ-line mutations. <i>Breast Cancer Research and Treatment</i> , 2008, 112, 343-349.	2.5	7
13	Disease family history and modification of breast cancer risk in common BRCA2 variants. <i>Oncology Reports</i> , 0, , .	2.6	4